

ATTORNEY'S DOCKET NO: 24702

U.S. DEPARTMENT OF COMMERCE, PATENT AND TRADEMARK OFFICE		DATE: <u>17</u> July 2001 (<u>17</u> .07.2001)
TRANSMITTAL LETTER TO THE UNITED STATES DESIGNATED/ELECTED OFFICE (DO/EO/US) CONCERNING A FILING UNDER 35 U.S.C. 371		U.S. APPL. NO. (Known) Not Yet Assigned 09/889348
INTERNATIONAL APPLICATION NO.: PCT/EP00/00324	INTERNATIONAL FILING DATE: 18 January 2000 (18.01.00)	PRIORITY DATE CLAIMED: 19 January 1999 (19 01.99)
TITLE OF INVENTION: RECOMBINANT SP-A FOR THE TREATMENT OR PREVENTION OF PULMONARY INFECTION AND INFLAMMATION		
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<p>Applicant hereby submits to the United States Designated/Elected Office (DO/EO/US) the following items and other information:</p> <ol style="list-style-type: none"> <input checked="" type="checkbox"/> This is a FIRST submission of items concerning a filing under 35 U.S.C. 371. <input type="checkbox"/> This is a SECOND or SUBSEQUENT submission of items concerning a filing under 35 U.S.C. 371. <input checked="" type="checkbox"/> This express request to begin national examination procedures (35 USC 371(f)) The submission must include items(5), (6), (9) and (21) indicated below. <input checked="" type="checkbox"/> A proper Demand for International Preliminary Examination was made by the 19th month from the earliest claimed priority date. <input checked="" type="checkbox"/> A copy of the International Application as filed (35 U.S.C. 371(c)(2)): <ol style="list-style-type: none"> <input checked="" type="checkbox"/> is transmitted herewith (required only if not transmitted by the International Bureau). <input type="checkbox"/> has been communicated by the International Bureau. <input type="checkbox"/> is not required, as the application was filed in the United States Receiving Office (RO/US) <input type="checkbox"/> A English translation of the International Application as filed (35 U.S.C. 371(c)(2)). <input checked="" type="checkbox"/> Amendments to the claims of the International Application under PCT Article 19 (35 U.S.C. 371(c)(3)) <ol style="list-style-type: none"> <input type="checkbox"/> are attached hereto (required only if not transmitted by the International Bureau). <input type="checkbox"/> have been communicated by the International Bureau. <input type="checkbox"/> have not been made; however, the time limit for making such amendments has NOT expired. <input checked="" type="checkbox"/> have not been made and will not be made. <input type="checkbox"/> An English language translation of the amendments to the claims under PCT Article 19 (35 U.S.C. 371(c)(3)). <input type="checkbox"/> An oath or declaration of the inventor(s) (35 U.S.C. 371(c)(4)). <input type="checkbox"/> A translation of the annexes to the International Preliminary Examination Report under PCT Article 36 (35 U.S.C. 371(c)(5)). <p>ITEMS 11 to 20 BELOW CONCERN OTHER DOCUMENT(S) OR INFORMATION INCLUDED:</p> <ol style="list-style-type: none"> <input type="checkbox"/> An Information Disclosure Statement under 37 CFR 1.97 and 1.98. <input type="checkbox"/> An assignment document for recording. A separate cover sheet in compliance with 37 CFR 3.28 and 3.31 is included. <input type="checkbox"/> A FIRST preliminary amendment. <input type="checkbox"/> A SECOND or SUBSEQUENT preliminary amendment <input type="checkbox"/> A substitute specification. <input type="checkbox"/> A change of power of attorney and/or address letter. <input type="checkbox"/> A computer-readable form of the sequence listing in accordance with PCT Rule 13ter2 and 35 USC 1821 - 1825 <input type="checkbox"/> A second copy of the published international application under 35 USC 154(d)(4) <input type="checkbox"/> A second copy of the English language translation of the international application under 35 USC 154(d)(4) <input checked="" type="checkbox"/> Other items or information: <p>TRANSMITTAL FORM; FEE CALCULATION; INTERNATIONAL PUBLICATION WO 00/43026; INTERNATIONAL PUBLICATION DATE 27 JULY 2000 CONSISTING OF 18 PAGES INCLUDING; 1 COVER SHEET CONTAINING THE ABSTRACT; 10 PAGES TEXTUAL SPECIFICATION, 2 PAGES OF 17 CLAIMS; 5 SHEETS DRAWINGS; PCT/RO/101 REQUEST; UNEXECUTED INVENTOR'S DECLARATION; PCT/IPEA/416 NOTIFICATION OF TRANSMITTAL OF INTERNATIONAL PRELIMINARY EXAMINATION REPORT; PCT/IPEA/409 INTERNATIONAL PRELIMINARY EXAMINATION REPORT; PCT/IB/304 NOTIFICATION CONCERNING SUBMISSION OR TRANSMITTAL OF PRIORITY DOCUMENT; PCT/IB/308 NOTICE INFORMING THE APPLICANT OF THE COMMUNICATION OF THE INTERNATIONAL APPLICATION TO THE DESIGNATED OFFICES; PCT/IB/332 INFORMATION CONCERNING ELECTED OFFICES NOTIFIED OF THEIR ELECTION.</p>		

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U.S. APPLICATION NO. 09/889348 <small>known entity assigned</small>	INTERNATIONAL APPLICATION NO. PCT/EP00/00324	DATE: <u>17</u> July 2001 (<u>17</u> .07.2001)
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17. <input checked="" type="checkbox"/> The following fees are submitted: Basic National Fee (37 CFR 1.492(a)(1)-(5): Search Report has been prepared by the EPO or JPO:.....\$860.00 International preliminary examination fee paid to USPTO (37 CFR 1.482).....\$690.00 No international preliminary examination fee paid to USPTO (37 CFR 1.482) but international search fee paid to USPTO (37 CFR 1.445(a)(2)).....\$710.00 Neither international preliminary examination fee (37 CFR 1.482) nor international search fee (37 CFR 1.445(a)(2)) paid to USPTO\$1000.00 International preliminary examination fee (37 CFR 1.482) and all claims satisfied provisions of PCT Article 33(2)-(4).....\$ 100.00 <div style="text-align: right;">ENTER APPROPRIATE BASIC FEE AMOUNT =</div>	<u>CALCULATIONS</u> \$ 860.00 \$ 860.00	<u>PTO USE ONLY</u>
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Surcharge of \$130.00 for furnishing the oath or declaration later than <u> </u> 20 <u> </u> 30 months from the earliest claimed priority date (37 CFR 1.492(e)).	\$	
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CLAIMS	NO. FILED	NO. EXTRA	RATE		
TOTAL	17 -20=	0	X \$ 18.00	\$	0.00
INDEPENDENT	4 - 3=	1	X \$ 80.00	\$	80.00
Multiple dependent claims(s) (if applicable)			+ \$260.00	\$	0.00
TOTAL OF ABOVE CALCULATIONS =				\$	940.00
Reduction by 1/2 for asserting small entity, if applicable. (Note 37 CFR 1.9, 1.27, 1.28).				\$	0.00
SUBTOTAL =				\$	940.00
Processing fee of \$130.00 for furnishing the English translation later than <u> </u> 20 30 months from the earliest claimed priority date (37 CFR 1.492(f)).				\$	0.00
TOTAL NATIONAL FEE =				\$	940.00
Fee for recording the enclosed assignment (37 CFR 1.21(h)). The assignment must be accompanied by an appropriate cover sheet (37 CFR 3.28, 3.31). \$40.00 per property +				\$	0.00
TOTAL FEES ENCLOSED =				\$	940.00
				Amount to be: refunded	\$
				charged	\$

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U.S. APPLICATION NO. (if known) not yet assigned 09/889348	INTERNATIONAL APPLICATION NO. PCT/EP00/00324	DATE: <u>17</u> July 2001 (<u>17</u> .07.2001)
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a. ☒ One check in the amount of \$940.00 to cover the above fees is enclosed.

b. ☐ Please charge my Deposit Account No. 14-0112 in the amount of \$_____ to cover the above fees. (A duplicate copy of this sheet is enclosed.)

c. ☒ The Commissioner is hereby authorized to charge any additional fees which may be required, or credit any overpayment to Deposit Account No. 14-0112.


d. Fees are to be charged to a credit card ☐ WARNING: Information on this form may become public ☐ Credit Card Information should not be included on this form. ☐ Provide credit card information and authorization on PTO-2038 ☐

NOTE: Where an appropriate time limit under 37 CFR 1.494 or 1.495 has not been met, a petition to revive (37 CFR 1.137(a) or (b)) must be filed to request that the application be restored to pending status.

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Rev. 02/98

**Recombinant SP-A for the treatment or prevention
of pulmonary infection and inflammation**

09/889348
JC18 Rec'd PCT/PTO 1 7 JUL 2001

Technical field

The present invention relates to the novel use of recombinant surfactant protein A for the production of a medicament for the treatment of pulmonary infection and inflammation.

Prior art

Pulmonary surfactant plays an important role in maintaining the structural integrity of the alveoli by reducing surface tension. The surfactant consists mostly of a complex mixture of phospholipids and genetically distinct proteins referred to as surfactant protein A, B, C and D (also designated as SP-A, SP-B, SP-C and SP-D). It is synthesized by alveolar type II pneumocytes and secreted as tightly packed lamellar bodies into the alveoli (King, R. J.: Pulmonary Surfactant, J. Appl. Physiol. **1982**, 51, 1-8).

SP-A is hypothesized to play a role in protecting the lung from bacterial, viral, and fungal infections (Thiel, S., and Reid, K.: Structures and functions associated with the group of mammalian lectins containing collagen-like sequences, FEBS Lett. **1989**, 250, 78). In addition, in vitro, it has been shown that SP-A binds to various micro-organisms, acts as opsonin, enhances killing of micro-organisms by macrophages, down regulates pro-inflammatory cytokines such as TNF- α induced by LPS or microbial pathogens (reviewed in: Molecular Basis of Disease, Pulmonary surfactant. Ed: L.M.G. van Golde, Biochimica et Biophysica Acta, **1998**, 1408, 77-364).

Recently, it was shown that mice lacking SP-A are susceptible to group B streptococcal infection (LeVine A. M. et al.: Surfactant protein A-deficient mice are susceptible to group B streptococcal infection; The Journal of Immunology, **1997**, 4336-4340) and that exogenous proteinosis SP-A enhanced bacterial clearance in SP-A deficient mice (LeVine A. M. et al.: Surfactant protein-A (SP-A) binds group B streptococcus (GBS), enhancing phagocytosis and clearance from lungs of SP-A deficient mice; Am J. Respir. Crit. Care Med. **1998**, Vol. 157, A 865). In addition, it was shown that baboons with bronchopulmonary dysplasia (BPD) and superimposed infection have decreased levels of SP-A present in the lungs (King, R. J., et al.: Surfactant protein-A deficiency in a primate model of pulmonary dysplasia, Am. J. Respir. Crit. Care Med. **1995**, 151(6), 1989-97 and Coalson, J. J.: Pathophysiologic, morphometric, and biochemical studies of the premature baboon with bronchopulmonary dysplasia, Am. Rev. Respir. Dis. **1992**, 145, 872-81). Furthermore, it was shown that SP-A is decreased in a number of diseases such as pneumonia, asthma, bronchiolitis, lung transplantations, cystic fibrosis, ARDS, smokers etc. as reviewed in M. Griese, Pulmonary surfactant in health and human lung diseases: state of the art. Eur. Respir. J. **1999**, 13, 1455-1476. It was also shown, that SP-A inhibits allergen induced histamine release as well as the proliferation of lymphocytes in cells isolated from allergen exposed asthmatics. (Wang, J.Y. et al., Inhibitory effect of pulmonary surfactant proteins A and D on allergen-induced lymphocyte proliferation and histamine release

in children with asthma. Am. J. Respir. Crit. Care Med., 1998, 158, 510-518; Mandan, T. et al., Lung surfactants proteins A and D can inhibit specific IgE binding to the allergens of *Aspergillus fumigatus* and block allergen-induced histamine release from human basophils. Clin. Exp. Immunol., 1997, 110, 241-249).

Summary of the Invention

The subject invention has several distinct aspects. One aspect is the use of a component which is at least substantially the same as recombinant surfactant protein A (rSP-A) for treating or preventing a pulmonary infection or inflammation. Another aspect is a medicament composition for treating or preventing a pulmonary infection and inflammation, and which comprises an active component which is at least substantially the same as recombinant surfactant protein A. A further aspect is a method of compounding such a medicament composition. A still further aspect comprises the concurrent use of surfactant protein D (SP-D) with an active component which is at least substantially the same as recombinant surfactant protein A in treating or preventing a pulmonary infection and inflammation, in compounding a medicament composition and in the resulting medicament composition itself. An additional aspect of the invention is an article of manufacture, comprising packaging material and rSP-A or a compound which is substantially the same as rSP-A (in a container) within the packaging material, and the packaging material including a label or instructions which indicate usefulness for the treatment or prevention of inflammation or microbial infection.

Brief Description of the Drawings

Figure 1 graphically illustrates that the clearance of GBS in the SP-A (-/-) mice was significantly enhanced at 6 and 24 hours when GBS was co-administered with 150 µg rSP-A.

Figure 2 graphically illustrates that GBS clearance by SP-A (-/-) mice is dose dependent and is increased by increased amount of rSP-A.

Figure 3 graphically illustrates the clearance of GBS in wild-type mice is significantly enhanced at 6 and 24 hours when GBS is co-administered with 150 µg rSP-A.

Figure 4 graphically illustrates that wild-type animals infected with GBS and treated 6 hours after infection with intratracheal rSP-A have increased clearance of GBS at 24 hours.

Figure 5 graphically illustrates that exogenous rSP-A reduces TNF-α content in lung homogenates from SP-A (-/-) mice challenged with GBS close to the level observed in SP-A (+/+) mice.

Details

Surprisingly it has now been found that recombinant surfactant-associated protein A (rSP-A) can be used in the treatment or prevention of pulmonary infection and inflammation and is equivalent or superior to the use of surfactant-associated protein A (SP-A) obtained from natural sources, for example that isolated from lavage fluid from healthy individuals or proteinosis patients. This must be regarded as particularly surprising as SP-A isolated from human lung lavage consists of a homogenous population of a flower bouquet like

hexameric structure, each unit of which consists of three SP-A polypeptide chains ($\alpha 1$), analogous to that described for the complement factor C1q. The fully assembled hexameric structure is thought to be essential for a functional molecule with respect to stimulating anti-microbial defense mechanisms or anti-inflammatory activity. In contrast to the naturally derived SP-A, surfactant-associated protein A produced by recombinant techniques consists of a variety of different oligomeric structures ranging from one single polypeptide chain ($\alpha 1$) to the fully assembled octadecameric form ($\gamma 6$) (Voss, T., et al.: Macromolecular organization of natural and recombinant lung surfactant protein SP 28-36; J. Mol. Biol. 1988, 201, 219-227; Voss et al.: Structural comparison of recombinant pulmonary surfactant protein SP-A derived from two human coding sequences: Implications for the chain composition of natural human SP-A, Am. J. Respir. Cell Mol. Biol., 1991, 4, 88-94).

In addition it was shown, that natural derived SP-A is glycosylated. However, although recombinant produced SP-A, depending on the system used for expression (mammalian, insect, or yeast cells), shows different glycosylation patterns, it shows anti-microbial or anti-inflammatory effects superior or equivalent to the natural SP-A.

As used herein microbial refers to bacterial, viral or fungal.

As used herein recombinant surfactant-associated protein A (hereinafter also referred to as rSP-A) refers to vertebrate, preferably mammalian, surfactant-associated protein A produced by recombinant techniques. Amino acid sequences and DNA sequences coding for mammalian surfactant-associated protein A are, for example, described in WO86/03408, WO88/05820 and USP 4,882,422. Recombinant surfactant-associated protein A further refers to derivatives of vertebrate, preferably mammalian, surfactant-associated protein A produced by recombinant techniques which differ from natural mammalian or other vertebrate surfactant-associated protein A by addition, deletion or substitution of amino acids as long as the proteins retain microbial clearance activity or anti-inflammatory activity. Such activity of recombinant surfactant-associated protein A can, for example, be determined in an assay according to the one herein-after described for group B streptococcus bacteria. In a preferred embodiment rSP-A refers to human surfactant-associated protein A produced by recombinant methods and having an amino acid sequence encoded by a DNA sequence contained in cDNA clones, pHS10-5, pHS10-4, PSAP-1A, PSAP-6A and a genomic clone pHS-15 or an allelic variation thereof. Two genes (designated as A1 and A2) coding for SP-A have been identified in the human genome (White, R. T. et al.: Nature 1985, 317, 361-363; Katyal, S. L. et al.: Am. J. Respir. Cell Mol. Biol., 1992, 6:446-452). The genomic clone pHS-15 coding for human SP-A has been described in WO86/03408, and clones containing cDNA sequences coding for SP-A have been described in WO88/05820 for pHS10-5 and pHS10-4, and by Floros et al., The Journal of Biological Chemistry, 1986, 261, 9029-33 and USP 4,882,422 for PSAP-1A and PSAP-6A). Recombinant SP-A may be obtained according to procedures known in the art. Methods for cloning and production of rSP-A are for example described in WO86/03408, WO88/05820, USP 4,659,805, USP 4,912,038, Voss, T., et al.: Macromolecular organization of natural and recombinant lung surfactant protein SP 28-36; J. Mol. Biol. 1988, 201, 219-227, and Voss et al.: Structural comparison of recombinant pulmonary surfactant protein SP-A derived from two human coding sequences: Implications for the chain composition of natural human SP-A, Am. J. Respir. Cell Mol. Biol., 1991, 4 88-94).

In addition, to express the cDNA sequences coding for SP-A (A1/A2) it is preferred to use either insect cells using the Baculovirus expression system (McCormack, F. et al., J. Biol. Chem., 1994, 269:5833-5841) or yeast, such as *Pichia pastoris*, in both cases with or without co-expression of the human prolyl 4-hydroxylase stabilizing the collagen helices by hydroxylating proline residues in the collagenous domain as demonstrated for the expression of collagen (Lamberg, Arja et al., J. Biol. Chem., 1996, 271:11988-

11995; Vuorela, A. et al., EMBO J., 1997, 16:6702-6712). For example rSP-A can be produced by cloning of the respective cDNAs into the EcoRI site of the Baculovirus expression vector pVL1392, subsequent generation of recombinant viruses and expression in SF21 cells using standard procedures. rSP-A may also be produced in yeast (for example *Pichia pastoris*) after cloning of the respective cDNAs into yeast expression vectors such as pPICZ A (Invitrogen, Leek, Netherlands) for the expression in *Pichia pastoris*.

In another embodiment of the invention recombinant SP-A obtainable by expression of a DNA sequence coding for SP-A in a suitable procaryotic expression system is used for the manufacture of a medicament for the prevention or treatment of pulmonary infection and inflammation.

In a further embodiment of the invention non-glycosylated rSP-A is used for the manufacture of a medicament for the prevention or treatment of pulmonary infection and inflammation.

A still further embodiment of the invention is an article of manufacture which comprises packaging material and a pharmaceutical agent within the packaging material wherein the pharmaceutical agent is at least substantially the same as rSP-A, and wherein the packaging material comprises a label or package insert which indicates that the pharmaceutical agent is useful for preventing or treating a pulmonary microbial infection or a pulmonary inflammation. The packaging material, label and package insert otherwise parallel or resemble what is generally regarded as standard packaging material, labels and package inserts for pharmaceuticals having related utilities.

Pharmacology

METHODS

Animal Husbandry

The murine SP-A gene locus was targeted by homologous recombination as previously described (Korfhagen; T.R. et al.: "Altered surfactant function and structure in SP-A gene targeted mice; PNAS, 1996, 93, 9594-9599). Lungs of SP-A (-/-) mice do not contain detectable SP-A mRNA or protein. To limit variability related to strain differences, 129 J wild type (+/+) and SP-A (-/-) mice of the same strain were studied. Animals were housed and studied under IACUC-approved protocols in the animal facility of the Children's Hospital Research Foundation, Cincinnati. Male and female mice of approximately 20 to 25 grams (35 to 42 days old) were used. (LeVine A. M. et al.: Surfactant protein A-deficient mice are susceptible to group B streptococcal infection; The Journal of Immunology, 1997, 158, 4336-4340).

Recombinant SP-A

Recombinant SP-A was produced as described by Voss et al. (Am. J. Respir. Cell Mol. Biol. 1991, 4, 88-94).

Preparation of Bacteria

A stock culture of group B streptococcus (GBS) was obtained from a clinical isolate from a newborn with systemic infection. Bacteria were suspended in sterile phosphate-buffered saline (PBS) containing 20 % glycerol and frozen in aliquots at -70°C . Bacteria from the same passage were used to minimize variations in virulence related to culture conditions. Before each experiment, an aliquot was thawed and plated on tryptic soy-5 % defibrinated sheep blood agar then inoculated into 4 ml of Todd-Hewitt broth (Difco Laboratories, Detroit, MI) and grown for 14 to 16 hours at 37°C with continuous shaking. The broth was centrifuged, and the bacteria were washed in PBS at pH 7.2 and resuspended in 4 ml of the buffer. In order to facilitate studies, a growth curve was generated so the bacterial concentration could be determined spectrophotometrically, which was confirmed by quantitative culture of the intratracheal inoculum.

Intratracheal Inoculation

Administration of GBS into the respiratory tract of the mice was performed by intratracheal inoculation of 10^4 or 10^5 cfu diluted in sterile normal saline (0.9 % NaCl). To deliver GBS in the presence of rSP-A, GBS was diluted in 0.9 % NaCl with 1 mM Ca^{2+} and appropriate amounts of rSP-A in a 37°C water bath for 30 minutes. Bacteria were delivered by intratracheal inoculation as previously described (LeVine A. M. et al.: Surfactant protein A-deficient mice are susceptible to group B streptococcal infection; *The Journal of Immunology*, 1997, 4336-4340).

Bacterial Clearance

Quantitative cultures of lung homogenates were performed 6 and 24 hours after inoculation of the animals with bacteria or bacteria together with SP-A, as previously described (LeVine A. M. et al.: Surfactant protein A-deficient mice are susceptible to group B streptococcal infection; *The Journal of Immunology*, 1997, 4336-4340). Bacterial clearance from the lungs of SP-A (-/-) mice was determined after intratracheal inoculation with GBS together with varying doses of rSP-A ranging from 25 μg , 50 μg , 75 μg , 100 μg to 150 μg . Wild-type animals were infected with GBS (10^6 cfu) and treated with rSP-A (150 μg) at the time of infection or 6 hours after infection.

Cytokine Production

Lung homogenates were centrifuged at $1,200 \times g$ and the supernatants were stored at -20°C . Tumor necrosis factor- α (TNF- α) levels were measured with ELISAs, using goat antimurine antibody (R&D Systems) directed against TNF- α . All plates were read on a microplate reader (Molecular Devices, Menlo Park, CA) and analyzed with the use of a computer-assisted analysis program (Softmax, Molecular Devices).

Statistical Methods

Since the distribution of the variable cfu/gram of lung was not normally distributed, a natural log transformation was used for all analyses. Analyses of variance (ANOVA) was performed to assess differences between the groups. Individual scores for each time point were compared using the median scores non parametric test. Findings were considered statistically significant at probability levels <0.05 .

RESULTS

Recombinant SP-A Increased Bacterial Clearance in SP-A (-/-) Mice

The clearance of GBS in the SP-A (-/-) mice was significantly enhanced at 6 and 24 hours when GBS was co-administered with 150 μ g rSP-A (Figure 1). Effects of rSP-A were comparable to that observed with SP-A isolated from proteinosis patients. GBS clearance by SP-A (-/-) mice was dose dependent and was increased by 50 μ g, 75 μ g, 100 μ g and 150 μ g of rSP-A (Figure 2).

Recombinant SP-A Increased Bacterial Clearance in Wild-type Mice

The clearance of GBS in wild-type mice was significantly enhanced at 6 and 24 hours when GBS was co-administered with 150 μ g rSP-A (Figure 3). Wild-type animals infected with GBS and treated 6 hours after infection with intratracheal rSP-A (150 μ g) had increased clearance of GBS at 24 hours (Figure 4).

Pulmonary clearance of intratracheally administered GBS was reduced in SP-A (-/-) mice compared to wild-type mice. Co-administration of exogenous recombinant SP-A with the bacteria significantly improved bacterial clearance demonstrating an immediate reversible defect in the SP-A (-/-) mouse. Enhanced pulmonary clearance of GBS with rSP-A treatment was dose dependent. Wild-type mice were effectively treated with rSP-A with increased clearance of GBS from the lungs.

In addition, it could be shown that exogenous rSP-A reduces TNF- α content in lung homogenates from SP-A (-/-) mice challenged with GBS close to the level observed in SP-A (+/+) mice (Figure 5).

Utility

On account of its microbial clearance and anti-inflammatory properties rSP-A is useful for the manufacture of medicaments for the prevention or treatment of pulmonary infection and inflammation. As used herein pulmonary infection refers to microbial pneumonias caused, for example, by viruses like *Respiratory Syncytial Virus*, *Adenovirus*, *Herpes simplex*, *Influenza A* and others or bacteria like, *Pseudomonas aeruginosa*, *Staphylococcus aureus*, *Haemophilus influenzae*, *Klebsiella pneumoniae*, *Group B Streptococci*, *Enterobacter* or *Streptococcus pneumoniae* and others, as well as fungi like *Aspergillus fumigatus*, *Pneumocystis carinii*, *Candida albicans* and others.

Furthermore pulmonary infection also refers to cases of reduced immunity for example the immunoparalytic phase occurring during sepsis and to immunodeficiency syndromes, whether congenital, spontaneously acquired, or iatrogenic. They are characterized by unusual susceptibility to infection and not infrequently to autoimmune disease and lymphoreticular malignancies. Patients with defects in humoral immunity have recurrent or chronic sinopulmonary infection, meningitis, and bacteremia, most commonly caused by pyogenic bacteria, such as *Haemophilus influenzae*, *Streptococcus pneumoniae*, and *Staphylococci*. These and other pyogenic organisms also cause frequent infections in individuals who have either neutropenia or a deficiency of the pivotal third component of complement (C3).

In connection with the present invention pulmonary inflammation refers, e.g., to bronchopulmonary dysplasia (BPD), and rSP-A therefore is also useful for the manufacture of medicaments for the prevention or treatment of bronchopulmonary dysplasia (BPD) or other disorders of SP-A deficiency. In particular BPD caused by artificial ventilation of premature babies may be mentioned in connection with the present invention. Additionally, inflammation also refers to pulmonary inflammation caused by artificial ventilation or cases of release of cytokines into the lung not primarily due to bacteria, viruses or fungi. Inflammation includes also diseases like asthma, CF (cystic fibrosis) and COPD (chronic obstructive pulmonary disease). It also includes acute lung injury up to the worst stages known as ARDS. In addition, inflammation also refers to inflammation induced by allergens.

The present invention also refers to the treatment of bacterial pulmonary infections with rSP-A or suitable forms thereof in combination or addition to antibiotics in the way that the infection induced inflammation is cured.

The invention furthermore relates to a method for the treatment of mammals, including humans, who are suffering from one of the above-mentioned illnesses. The method is characterized in that a therapeutically active and pharmacologically tolerable amount of rSP-A is administered to the mammal in need thereof.

In connection with the novel use of rSP-A according to the invention medicaments are prepared by procedures familiar to those skilled in the art. To do this rSP-A is either employed as such or preferably in combination with suitable pharmaceutical auxiliaries, e.g., as suspensions, solutions or in powder form, the rSP-A content advantageously being from 0.1 to 90 % (wt/wt), preferably 0.1 to 15 % (wt/wt). The rSP-A can be administered either alone or with auxiliaries. The auxiliaries which are suitable for the desired pharmaceutical formulations are familiar to the person skilled in the art on account of his expert knowledge. Pharmaceutical formulations which can be used according to the present invention and containing rSP-A in combination with synthetic or natural lipids are, for example, disclosed in USP 4,659,805. Surprisingly, and in contrast to pharmaceutical formulations comprising rSP-A disclosed in the state of the art, lipid free formulations comprising rSP-A as active ingredient can also be used for the treatment of pulmonary infection and inflammation. A further aspect the invention thus relates to lipid free medicaments comprising rSP-A.

In particular such lipid free medicaments comprising rSP-A contain as auxiliary from 0.01 to 0.1 % of a calcium salt, preferably calcium chloride. An exemplary liquid, lipid free medicament can be manufactured

by dissolving 100 to 1,000 mg of rSP-A and 11.1 mg of calcium chloride in 100 ml of a sterile 0.9 % aqueous sodium chloride solution.

In a preferred embodiment the medicaments are made available in liquid form for intratracheal or intrabronchial administration by instillation or nebulization or in powder form for administration by inhalation.

In connection with the novel use of rSP-A according to the invention medicaments are administered, for example, 2 to 3 times daily for from 1 to 7 days. For example medicaments comprising 100µg/kg to 10mg/kg (of body weight) of rSP-A are administered by inhalation or intratracheally or intrabronchially.

In a further aspect of the invention rSP-A is administered in combination with surfactant proteins SP-B, SP-C and/or SP-D or their modified derivatives for the treatment or prevention of pulmonary infection and inflammation. In particular the co-administration of rSP-A with SP-D is preferred in connection with the invention. The amino acid sequences of said surfactant proteins, their isolation or preparation by genetic engineering are known (e.g. from WO86/03408, EP-A-0 251 449, WO89/04326, WO87/06943, WO88/03170, EP-A-0 368 823, EP-A-0 348 967, WO91/18015, WO95/32992, EP-A-0 593 094, WO92/22315 and Crouch et al.: Recombinant Pulmonary Surfactant Protein D, The Journal of Histological Chemistry, 1994, 269, 15808-15813 and references cited therein).

Description of figures

Figure 1: Enhanced clearance of GBS from the lung in SP-A (-/-) mice. 10^4 cfu GBS were inoculated with or without 150 µg of rSP-A, and colony counts were performed after 6 and 24 hours as previously described. SP-A (-/-) without rSP-A (solid bars) and with rSP-A (hatched bars). Data are Means \pm SEM (standard error of means) values.

Figure 2: Enhanced clearance of GBS by exogenous rSP-A is dose dependent. SP-A (-/-) mice were inoculated with 10^4 cfu GBS in the presence of either 0 (N.T.), 25, 50, 75, 100 or 150 µg of rSP-A, and colony counts were performed 6 hours after intratracheal instillation as previously described. Data are Means \pm SEM values.

Figure 3: Enhanced clearance of GBS from the lung in SP-A (+/+) mice (wt). 10^6 cfu GBS were inoculated with or without 150 µg of rSP-A and colony counts were performed 6 and 24 hours after intratracheal instillation as previously described. SP-A (+/+) without rSP-A (hatched bars) and with rSP-A (solid bars). Data are Means \pm SEM values.

Figure 4: Rescue of GBS infection in SP-A(wt) mice. SP-A (+/+) mice were intratracheally inoculated with 10^6 cfu GBS in the absence of exogenous rSP-A. 6 hours after infection, 150 µg of rSP-A were intratracheally administered. 24 hours after rSP-A administration colony counts in lungs were performed. Colony counts were dramatically reduced in lungs from animals treated with rSP-A (solid bars) compared to untreated animals (hatched bars). Data are Means \pm SEM values.

The invention and its advantages are readily understood from the preceding description. Various changes may be made in the processes and compositions without departing from the spirit and scope of the invention or sacrificing its material advantages, the processes and products hereinbefore described being merely illustrative of preferred embodiments of the invention.

WHAT IS CLAIMED IS:

1. In a method for compounding a medicament composition for preventing or treating a pulmonary infection or inflammation and comprising suitable carrier and a pharmaceutically-acceptable active component, the improvement wherein the active component is at least substantially the same as recombinant surfactant protein A (rSP-A).
2. A method of claim 1 wherein rSP-A is recombinant SP-A which is the same as that obtainable by expression of a genomic sequence coding for SP-A in a suitable expression system.
3. A method of claim 1 wherein rSP-A is recombinant SP-A which is the same as that obtainable by expression of a cDNA coding for SP-A in a suitable expression system.
4. A method of claim 1 wherein the medicament composition further comprises surfactant protein D (SP-D).
5. A lipid-free medicament composition which is useful for treating or preventing a pulmonary infection or inflammation and which comprises an effective amount of pharmaceutically-acceptable active component and suitable carrier therefor, wherein the active component is at least substantially the same as rSP-A.
6. A medicament composition of claim 5 for treating or preventing a pulmonary infection.
7. A medicament composition of claim 6 wherein the pulmonary infection is bacterial, viral or fungal pneumonia.
8. A medicament composition of claim 5 for treating or preventing a pulmonary inflammation.
9. A medicament composition of claim 8 wherein the pulmonary inflammation is bronchopulmonary dysplasia.
10. A medicament composition of claim 5 wherein the rSP-A is at least substantially the same as recombinant SP-A obtained by expression of a genomic sequence coding for SP-A in a suitable expression system.
11. A medicament composition of claim 5 wherein the rSP-A is at least substantially the same as recombinant SP-A obtained by expression of a cDNA coding for SP-A in a suitable expression system.

12. A medicament composition of claim 5 wherein the active component further comprises surfactant protein D (SP-D).
13. In a method for preventing or treating a pulmonary infection or inflammation in a subject prone to or afflicted with such condition, which method comprises administering an effective amount of pharmaceutically-acceptable active component to the subject, the improvement wherein the active component comprises an essential ingredient which is at least substantially the same as recombinant surfactant protein A (rSP-A).
14. A method of claim 13 wherein the essential ingredient is rSP-A.
15. A method of claim 13 wherein the active component further comprises surfactant protein D (SP-D).
16. An article of manufacture comprising packaging material and a pharmaceutical composition contained within the packaging material, wherein the pharmaceutical composition comprises an active agent which is at least substantially the same as recombinant surfactant protein A, and wherein the packaging material comprises a label or package insert which indicates that the active agent is useful for preventing or treating a pulmonary microbial infection or inflammation.
17. An article of manufacture of claim 16 wherein the active agent is rSP-A.

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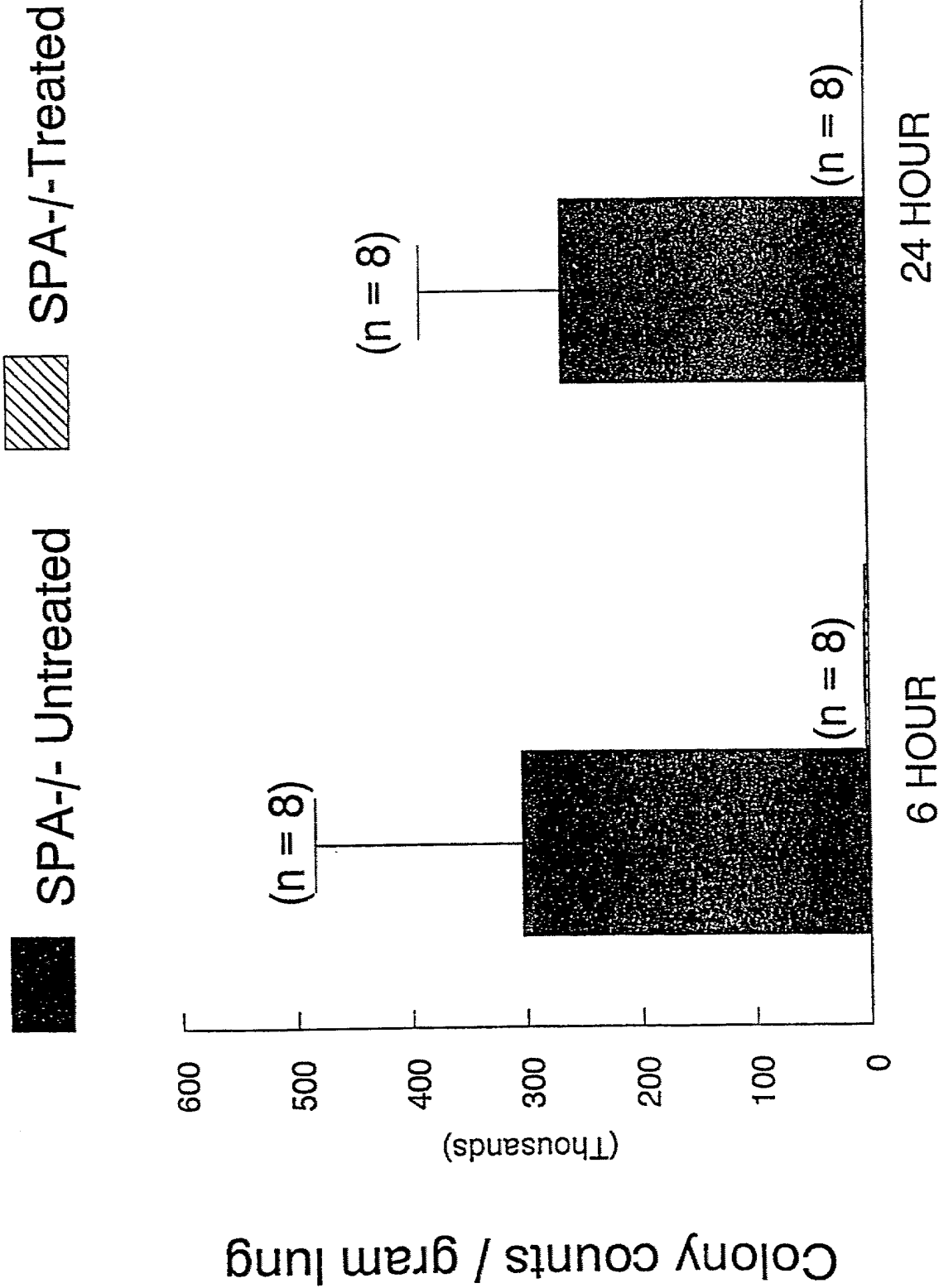


Figure 1

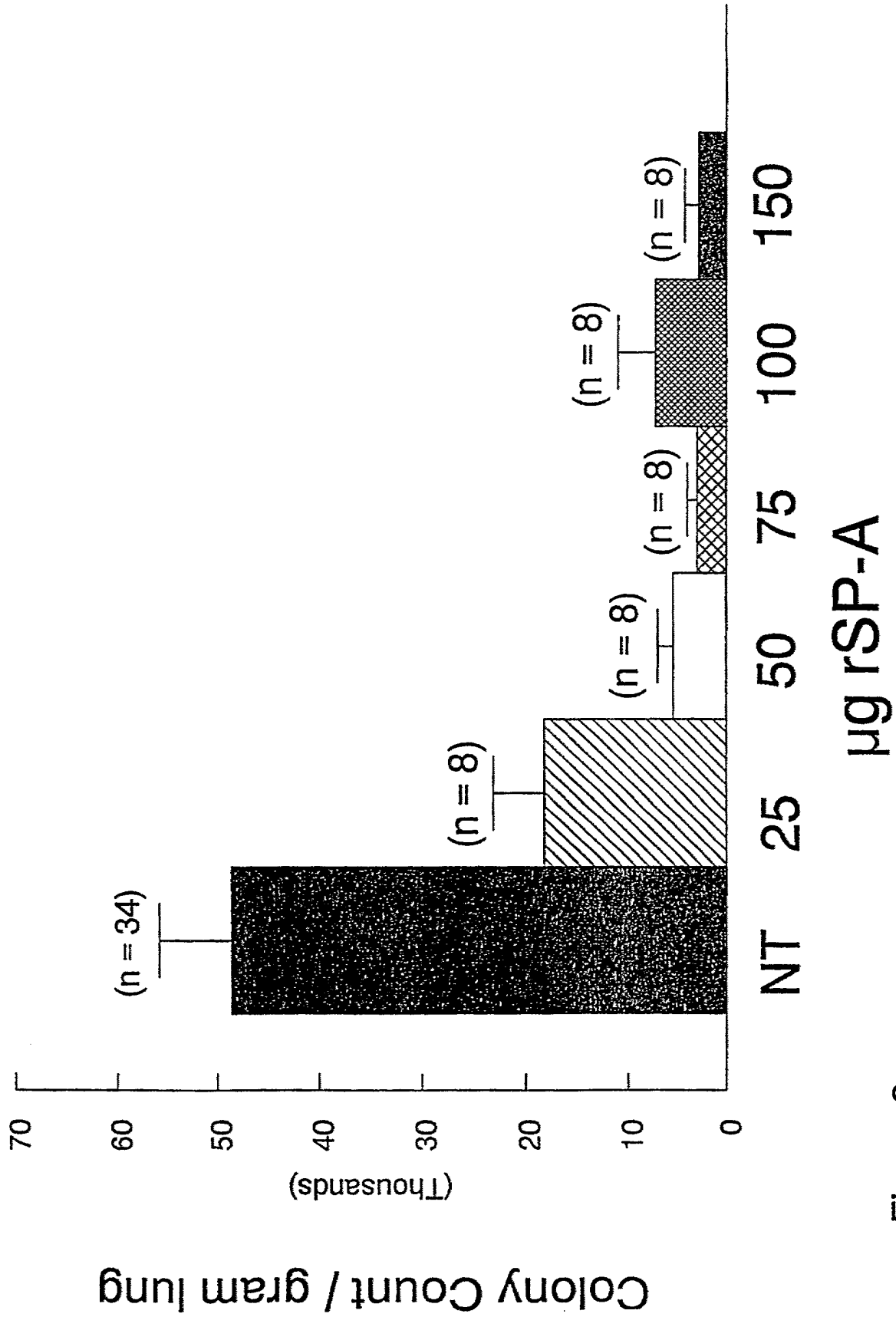


Figure 2

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3/5

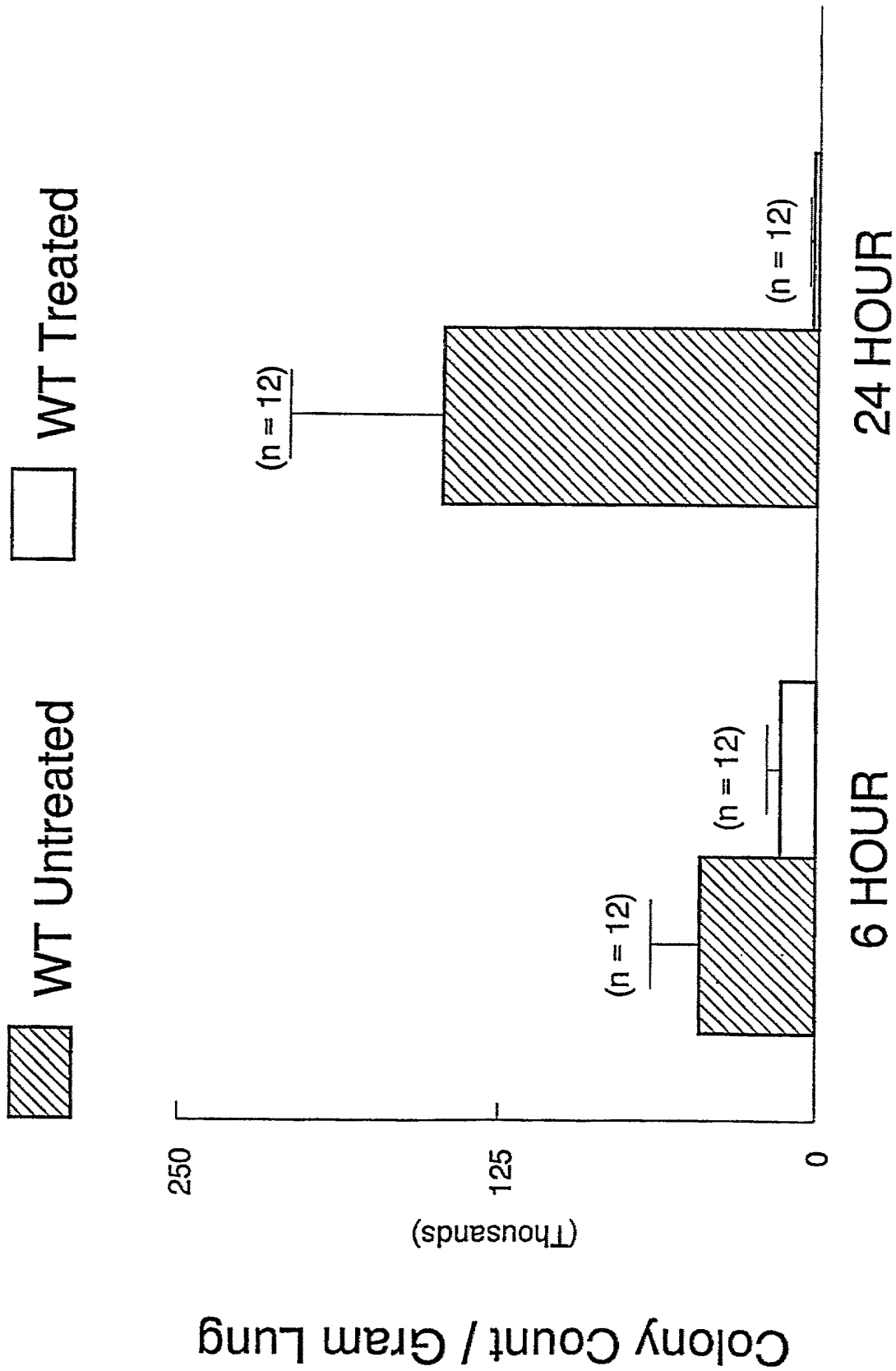


Figure 3

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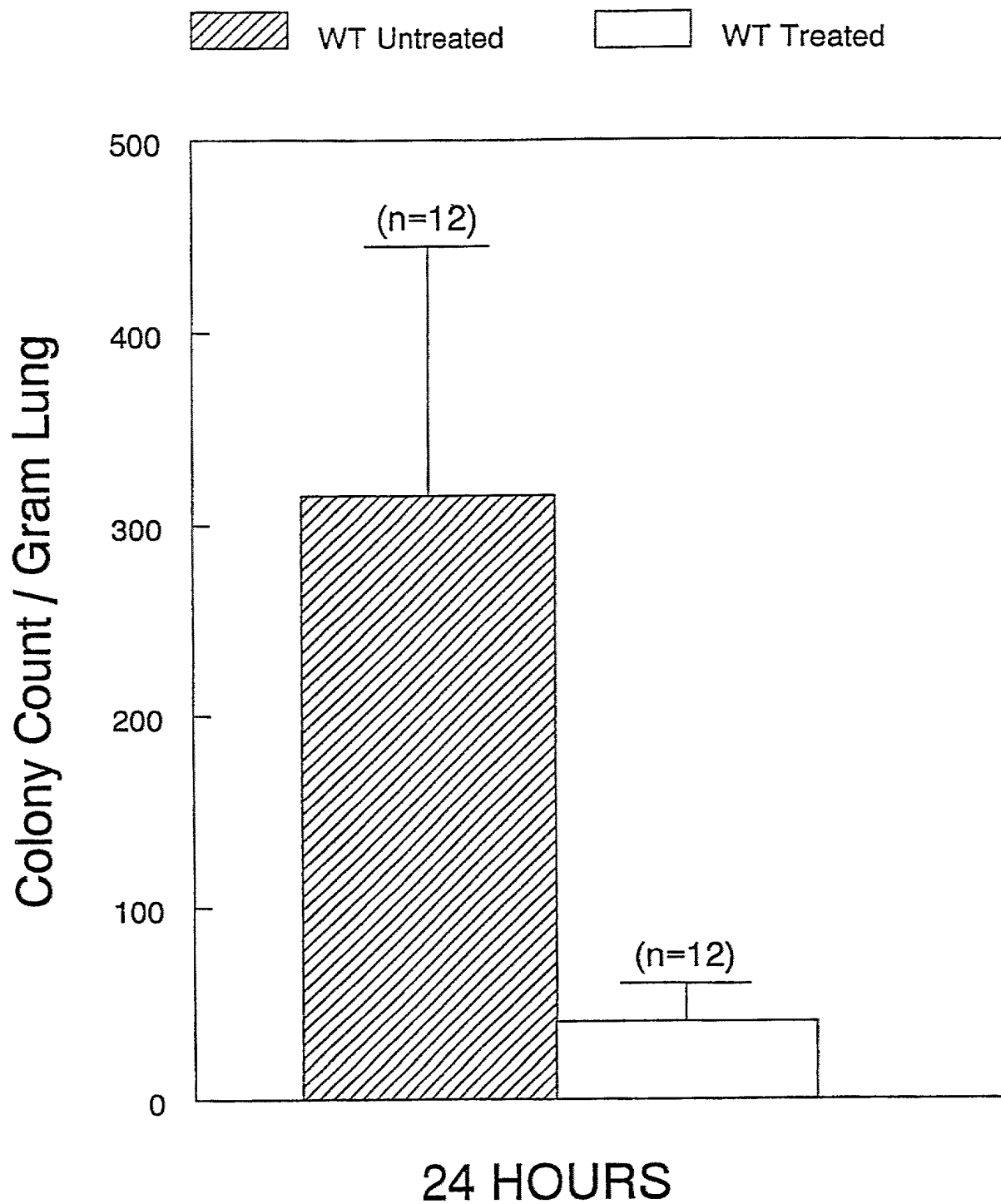


Figure 4

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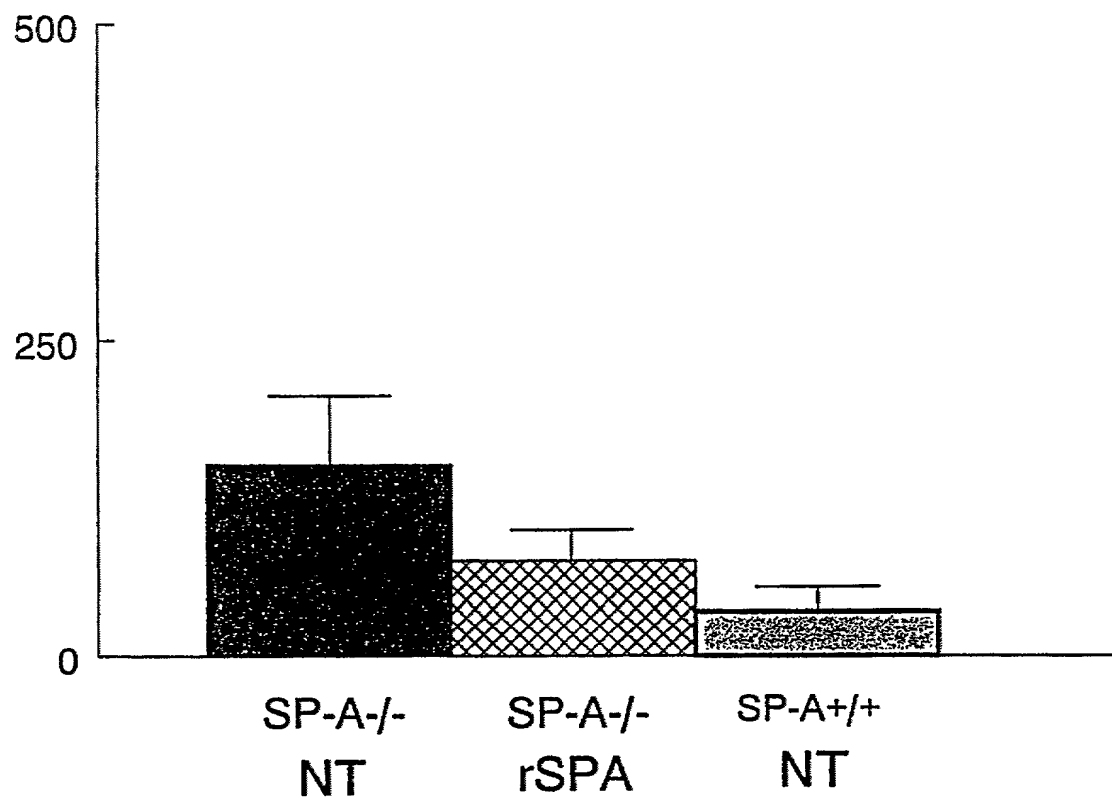


Figure 5

DECLARATION FOR PATENT APPLICATION

Attorney Docket: 24702

Page 1 of 2

As a below-named inventor(s), I/we hereby declare that:

My/Our residence(s), post office address(es) and citizenship(s) is/are as stated below next to my/our name(s).

I/We believe I/we am/are the original inventor, first and sole (if only one name is listed below) or the original, first and joint inventors (if plural names are listed below) of the subject matter which is claimed, and for which a patent is sought on the invention entitled:

"Recombinant SP-A for the treatment or prevention of pulmonary infection and inflammation"
the specification of which: (check one)

☐ is attached hereto.

☒ was filed on 18.01.2000, as Serial No. PCT/EP00/00324,

and was amended on _____ (if applicable).

We hereby state that we have reviewed and understand the contents of the above-identified specification, including the claims, as amended by any amendment referred to above.

We acknowledge the duty to disclose information which is material to the patentability of this application as defined by 37 CFR § 1.56.

We hereby claim foreign priority benefits under 35 U.S.C. § 119 of any foreign application(s) for patent or inventor's certificate listed below, and have also identified below any foreign application for patent or inventor's certificate having a filing date before that of the application on which priority is claimed:

Prior Foreign Applications:

			Priority Claimed
<u>60/155268</u>	<u>USA</u>	<u>19/01/1999</u>	<input checked="" type="checkbox"/> <input type="checkbox"/>
(Application No.)	(Country)	(Day/Month/Year Filed)	Yes No
 (Application No.)	 (Country)	 (Day/Month/Year Filed)	<input type="checkbox"/> <input type="checkbox"/>
 (Application No.)	 (Country)	 (Day/Month/Year Filed)	<input type="checkbox"/> <input type="checkbox"/>
 (Application No.)	 (Country)	 (Day/Month/Year Filed)	<input type="checkbox"/> <input type="checkbox"/>

We hereby appoint Gary M. Nath, Reg. No. 26,965; Harold L. Novick, Reg. No. 26,011; Todd L. Juneau, Reg. No. 40,669; Lee C. Heiman, Reg. No. 41,827; Jerald L. Meyer, Reg. No. 41,194; Joshua B. Goldberg, Reg. No. 44,126; David R. Murphy, Reg. No. 22,751; Paul A. Sacher, Reg. No. 43,418; Deborah H. Yellin, 45,904; Nahied K. Usman, Reg. 47,148; and Roger Hahn, Reg. No. 46,376; as my attorneys to prosecute this application and transact all business in the U.S. Patent and Trademark Office connected therewith.

Direct Telephone Calls to:

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020529

PATENT TRADEMARK OFFICE

Send Correspondence to:

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We hereby claim the benefit under 35 U.S.C. § 120 of any United States application(s) listed below and, insofar as the subject matter of each of the claims of this application is not disclosed in the prior United States application in the manner provided by 35 U.S.C. § 112, first paragraph, I/we acknowledge the duty to disclose material information as defined in 37 CFR § 1.56 which occurred between the filing date of the prior application and the national or PCT international filing date of this application:

(U.S. Application Serial No.) (U.S. Filing Date) (Status--patented, pending, abandoned)



B650USPCT01

DECLARATION FOR PATENT APPLICATION

Attorney Docket: 24702
Page 2 of 2

We hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements are made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under 18 U.S.C. ' 1001 and that such willful false statements may jeopardize the validity of the application or any patent issued thereon.

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Inventor's Signature _____ Date: _____
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Post Office Address: 5565 Salem Road, CINCINNATI, OH 45230, USA

Full name of third inventor: LEVINE, Ann Marie

Inventor's Signature _____ Date: _____
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Full name of fourth inventor: KORFHAGEN, Thomas R.

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Full name of fifth inventor: _____

Inventor's Signature _____ Date: _____
Residence: _____
Country of Citizenship: _____
Post Office Address: _____

Full name of sixth inventor: _____

Inventor's Signature _____ Date: _____
Residence: _____
Country of Citizenship: _____
Post Office Address: _____

DECLARATION FOR PATENT APPLICATION

Attorney Docket: 24702

Page 2 of 2

We hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements are made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under 18 U.S.C. § 1001 and that such willful false statements may jeopardize the validity of the application or any patent issued thereon.

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Inventor's Signature *Jeffrey Whitsett* Date: 8/6/01

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Inventor's Signature *Thomas Korfhagen* Date: 8/9/01

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Inventor's Signature _____ Date: _____

Residence: _____

Country of Citizenship: _____

Post Office Address: _____

Full name of sixth inventor: _____

Inventor's Signature _____ Date: _____

Residence: _____

Country of Citizenship: _____

Post Office Address: _____